09/830744

L Number	Hits	Search Text	DB	Time stamp
1	54180		USPAT;	2002/09/23 13:45
			US-PGPUB;	
]	1		EPO; JPO; DERWENT;	
1			IBM TDB	
2	8944	polysaccharide and cross-link\$	USPAT;	2002/09/23 13:46
-		polyocomical and the same	US-PGPUB;	
			EPO; JPO;	
		•	DERWENT;	
	1050	t 1 land and annual limbely and	IBM_TDB USPAT;	2002/09/23 13:47
3	1052	(polysaccharide and cross-link\$) and polyamine	US-PGPUB;	2002/03/23 13.47
		poryamine	EPO; JPO;	
			DERWENT;	
			IBM_TDB	
4	328		USPAT;	2002/09/23 13:47
		polyamine) and carboxy	US-PGPUB; EPO; JPO;	
			DERWENT;	
		·	IBM TDB	
5	94	(((polysaccharide and cross-link\$) and	USPAT;	2002/09/23 13:49
]		polyamine) and carboxy) and activate	US-PGPUB;	
			EPO; JPO;	
			DERWENT; IBM TDB	
6	6568	polysaccharide and carboxy	USPAT;	2002/09/23 13:48
] "	0308	polysacchallac and calbony	US-PGPUB;	
			EPO; JPO;	
			DERWENT;	
			IBM_TDB	0000/00/03 13:40
7	2134	(polysaccharide and carboxy) and cross-link\$	USPAT; US-PGPUB;	2002/09/23 13:48
		Closs-links	EPO; JPO;	
			DERWENT;	
1 .			IBM_TDB	
8	476	((polysaccharide and carboxy) and	USPAT;	2002/09/23 13:49
		cross-link\$) and (diamine or triamine)	US-PGPUB; EPO; JPO;	
			DERWENT;	
			IBM TDB	
9	0	(((polysaccharide and carboxy) and	USPAT;	2002/09/23 13:50
		cross-link\$) and (diamine or triamine))	US-PGPUB;	
		and hyaluronicl4 and activate	EPO; JPO;	
			DERWENT; IBM TDB	
10	80	(((polysaccharide and carboxy) and	USPAT;	2002/09/23 13:50
		cross-link\$) and (diamine or triamine))	US-PGPUB;	
		and hyaluronic	EPO; JPO;	
1			DERWENT;	
111	67	(((polysaccharide and carboxy) and	IBM_TDB USPAT;	2002/09/23 14:18
**	6/	cross-link\$) and (diamine or triamine))	US-PGPUB;	
		and chitin	EPO; JPO;	
]			DERWENT;	
1		536/01	IBM_TDB	0000/00/00
12	535	536/21	USPAT;	2002/09/23 14:18
			US-PGPUB; EPO; JPO;	
			DERWENT;	
			IBM_TDB	
13	94	536/21 and cross-link\$	USPAT;	2002/09/23 14:18
			US-PGPUB;]
1	-		EPO; JPO; DERWENT;]
			IBM TDB	
14	27	(536/21 and cross-link\$) and hyaluronic	USPAT;	2002/09/23 14:18
		•	US-PGPUB;	
			EPO; JPO;	
			DERWENT; IBM TDB	
	l		TDM TOB	<u> </u>

15	6	((536/21 and cross-link\$) and hyaluronic)	USPAT;	2002/09/23 14:22
1	!	and (diamine or polyamine)	US-PGPUB;	
	1		EPO; JPO;	į
			DERWENT;	
			IBM_TDB	[
16	2833	514/54	USPAT;	2002/09/23 14:23
1			US-PGPUB;	
			EPO; JPO;	[
	i		DERWENT;	1
			IBM_TDB	
17	522	514/54 and cross-link\$	USPAT;	2002/09/23 14:23
		·	US-PGPUB;	
1 .			EPO; JPO;	
ļ			DERWENT;	
			IBM_TDB	
18	192	(514/54 and cross-link\$) and hyaluronic	USPAT;	2002/09/23 14:23
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			EPO; JPO;	· i
1			DERWENT;	
			IBM_TDB	
19	24	((514/54 and cross-link\$) and hyaluronic)	USPAT;	2002/09/23 14:23
		and (diamine or polyamine)	US-PGPUB;	
1			EPO; JPO;	
			DERWENT;	
			IBM TDB	

L Number	Hits	Search Text	DB	Time stamp
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-	3,100	F4	US-PGPUB;	
			EPO; JPO;	
		·	DERWENT;	
1 1			IBM TDB	}
2	8944	polysaccharide and cross-link\$	USPAT;	2002/09/23 13:46
]			US-PGPUB;	
1			EPO; JPO;	
			DERWENT;	
[·	IBM TDB	1
3	1052	(polysaccharide and cross-link\$) and	USPĀT;	2002/09/23 13:47
		polyamine	US-PGPUB;	1
			EPO; JPO;	
			DERWENT;	
]			IBM_TDB	İ
4	328		USPAT;	2002/09/23 13:47
i ,		polyamine) and carboxy	US-PGPUB;	
			EPO; JPO;	
1		·	DERWENT;	
l i			IBM_TDB	i .
5	94		USPAT;	2002/09/23 13:49
		polyamine) and carboxy) and activate	US-PGPUB;	
			EPO; JPO;	
			DERWENT;	
			IBM_TDB	
6	6568	polysaccharide and carboxy	USPAT;	2002/09/23 13:48
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			EPO; JPO;	1
			DERWENT;	
			IBM_TDB	
7	2134	1 14 1 4 1 1	USPAT;	2002/09/23 13:48
		cross-link\$	US-PGPUB;	
			EPO; JPO;	
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			IBM_TDB	
8	476	((polysaccharide and carboxy) and	USPAT;	2002/09/23 13:49
		cross-link\$) and (diamine or triamine)	US-PGPUB;	
			EPO; JPO;	
			DERWENT;	l .
9	ا ه	///palwasahawida and asahawul and	IBM_TDB USPAT;	2002/09/23 13:50
,	١	<pre>(((polysaccharide and carboxy) and cross-link\$) and (diamine or triamine))</pre>	US-PGPUB;	2002/09/23 13:50
		and hyaluronicl4 and activate	EPO; JPO;	
ļ	l	and hyaluronicia and accivace	DERWENT;	
. [i	•	IBM TDB	1
10	80	(((polysaccharide and carboxy) and	USPAT;	2002/09/23 13:50
	30	cross-link\$) and (diamine or triamine))	US-PGPUB;	2002/03/23 13:30
İ	i	and hyaluronic	EPO; JPO;	
1	ļ		DERWENT;	
			IBM TOB	
11	67	(((polysaccharide and carboxy) and	USPAT;	2002/09/23 14:18
	٠. ا	cross-link\$) and (diamine or triamine))	US-PGPUB;	= 332, 33, 23 13.10
ļ		and chitin	EPO; JPO;	
			DERWENT;	
. !			IBM TDB	
12	535	536/21	USPAT;	2002/09/23 14:18
			US-PGPUB;	
			EPO; JPO;	
			DERWENT;	
		,	IBM TDB	
13	94	536/21 and cross-link\$	USPAT;	2002/09/23 14:18
	ļ		US-PGPUB;	
	ļ		EPO; JPO;]
	1		DERWENT;	
i	ĺ		IBM TDB	
14	27	(536/21 and cross-link\$) and hyaluronic	USPAT;	2002/09/23 14:18
	ŀ	•	US-PGPUB;	
	ĺ		EPO; JPO;	
1	l		DERWENT;	
	ļ		IBM TDB	1

		Tirescion and annual limbor and burlumonia	LUCDATE.	1 2002/09/23 14:22
15	6	((536/21 and cross-link\$) and hyaluronic) and (diamine or polyamine)	USPAT; US-PGPUB;	2002/03/23 14.22
		and (diamine of polyamine)	EPO; JPO;	
1]	DERWENT;	
			IBM_TDB	
16	2833	514/54	USPAT;	2002/09/23 14:23
			US-PGPUB;	}
			EPO; JPO;	
	1		DERWENT; IBM TDB	
17	522	514/54 and cross-link\$	USPĀT;	2002/09/23 14:23
1 '	322	314754 und 51555 111114	US-PGPUB;	1
}			EPO; JPO;	1 *
			DERWENT;	1
	1		IBM_TDB	2000/00/03 14:03
18	192	(514/54 and cross-link\$) and hyaluronic	USPAT; US-PGPUB;	2002/09/23 14:23
1	İ		EPO; JPO;	
1			DERWENT;	
	1		IBM TDB	
19	24	((514/54 and cross-link\$) and hyaluronic)	USPĀT;	2002/09/23 14:49
	İ	and (diamine or polyamine)	US-PGPUB;	
ĺ	1	·	EPO; JPO;	
ĺ			DERWENT; IBM TDB	
20	0	(((514/54 and cross-link\$) and hyaluronic)	USPAT;	2002/09/23 14:49
20	, v	and (diamine or polyamine)) and complex?	US-PGPUB;	2002/03/20 21.13
			EPO; JPO;	
			DERWENT;	
			IBM_TDB	
21	130	((514/54 and cross-link\$) and hyaluronic)	USPAT;	2002/09/23 14:50
		and comple?	US-PGPUB; EPO; JPO;	
			DERWENT;	
			IBM TDB	
22	57	(((514/54 and cross-link\$) and hyaluronic)	USPĀT;	2002/09/23 14:52
		and comple?) and (copper or iron)	US-PGPUB;	
			EPO; JPO;	
	•		DERWENT; IBM TDB	
23	76	((((polysaccharide and carboxy) and	USPAT;	2002/09/23 14:58
		cross-link\$) and (diamine or triamine))	US-PGPUB;	
		and hyaluronic) and (copper or iron or	EPO; JPO;	
		metal or ion)	DERWENT;	
24	5	(((514/54 and cross-link\$) and hyaluronic)	IBM_TDB USPAT;	2002/09/23 14:58
23	,	and comple?) and salified	US-PGPUB;	2002/03/23 14.30
		·	EPO; JPO;	
		·	DERWENT;	
0.5	_		IBM_TDB	
25	3	((((514/54 and cross-link\$) and	USPAT;	2002/09/23 15:33
]	hyaluronic) and comple?) and salified) and (copper or iron or zinc)	US-PGPUB; EPO; JPO;	
		(copper or from or affic)	DERWENT;	
			IBM TDB	
26	22	(((514/54 and cross-link\$) and hyaluronic)	USPAT;	2002/09/23 15:00
		and (diamine or polyamine)) and sulfat?	US-PGPUB;	
	ĺ		EPO; JPO;	
			DERWENT; IBM TDB	
27	o	((((514/54 and cross-link\$) and	USPAT;	2002/09/23 15:00
		hyaluronic) and (diamine or polyamine))	US-PGPUB;	
į		and sulfat?) and trioxide	EPO; JPO;]
			DERWENT;	
28	30051	((((514/54 and cross-link\$) and	IBM_TDB	2002/00/22 15:01
-0	30331	hyaluronic) and (diamine or polyamine))	USPAT; US-PGPUB;	2002/09/23 15:01
		and sulfat?) and sulfur trioxide	EPO; JPO;	
			DERWENT;	
			IBM TDB	

29	2	((((514/54 and cross-link\$) and	USPAT;	2002/09/23 15:04
29		hyaluronic) and (diamine or polyamine))	US-PGPUB;	
		and sulfat?) and sulfation	EPO; JPO;	
		and Sallaci, and Sallacion	DERWENT;	
			IBM TDB	
30	ا ا	((((polysaccharide and carboxy) and	USPAT;	2002/09/23 15:04
30	ľ	cross-link\$) and (diamine or triamine))	US-PGPUB;	
		and hyaluronic) and sulfation	EPO; JPO;	
		and nyarazonzo, and outliness	DERWENT;	
		,	IBM TDB	
31	5	((536/21 and cross-link\$) and hyaluronic)	USPAT;	2002/09/23 15:06
J	1	and sulfation	US-PGPUB;	
			EPO; JPO;	1
			DERWENT;	
			IBM_TDB	·
32	3	(((536/21 and cross-link\$) and hyaluronic)	USPAT;	2002/09/23 15:06
		and sulfation) and pyridine	US-PGPUB;	
			EPO; JPO;	
			DERWENT;	
			IBM_TDB	
33	1	((514/54 and cross-link\$) and hyaluronic)	USPAT;	2002/09/23 15:34
		and complex?	US-PGPUB;	
			EPO; JPO;	
			DERWENT;	
	•		IBM_TDB	0000 (00 (00 15 06
34	1		USPAT;	2002/09/23 15:36
		and complex?) and (copper or zinc or iron)	US-PGPUB;	
			EPO; JPO;	
		•	DERWENT;	
~-		////F34/F4 and among link() and	IBM_TDB USPAT;	2002/09/23 15:36
35	0	(((())))	US-PGPUB;	2002/09/23 15:36
		hyaluronic) and complex?) and (copper or	EPO; JPO;	
		zinc or iron)) and cross-link?	DERWENT;	
			IBM TOB	
			TOM_TOD	i

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=> file polymers
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SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

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FILE 'USPAT2' ENTERED AT 14:31:14 ON 23 SEP 2002
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=> s polysaccharide
        218150 POLYSACCHARIDE
=> s ll and carboxy
          7277 L1 AND CARBOXY
=> s 12 and activat?
          3796 L2 AND ACTIVAT?
=> s 13 and cross-link
           317 L3 AND CROSS-LINK
=> s 14 and (diamine or polyamine)
L5
            76 L4 AND (DIAMINE OR POLYAMINE)
=> s 15 and hyaluroni
             0 L5 AND HYALURONI
L6
=> s 15 and hyaluronic
            24 L5 AND HYALURONIC
=> dis 17 1-24 bib abs
     ANSWER 1 OF 24 USPATFULL
1.7
       2002:235521 USPATFULL
AN
ΤI
       Process for ex vivo formation of mammalian bone and uses thereof
TN
       Kale, Sujata, Boston, MA, UNITED STATES
       Long, Michael W., Northville, MI, UNITED STATES
                                20020912
PT
       US 2002127711
                          A1
ΑI
       US 2000-753043
                           A1
                                20001227 (9)
DT
       Utility
FS
       APPLICATION
       Steven L. Highlander, Fulbright & Jaworski L.L.P.,, 600 Congress Avenue
LREP
       Suite 2400, Austin, TX, 78701
CLMN
       Number of Claims: 38
ECL
       Exemplary Claim: 1
DRWN
       10 Drawing Page(s)
LN.CNT 3032
       The present invention concerns methods for the ex vivo formation of
       mammalian bone and subsequent uses of the bone. A critical and
       distinguishing feature of the present invention are defined tissue
       culture conditions and factors resulting in the formation of bone cell
       spheroids. The invention also provides for methods of implanting into
       subjects the ex vivo formed bone. Also described are methods for
       genetically altering the bone cell spheroids to affect bone formation,
       identification of candidate modulators of bone formation, and
       identification of genes involved in bone formation.
L7
     ANSWER 2 OF 24 USPATFULL
       2002:224605 USPATFULL
AN
TI
       Lipid soluble steroid prodrugs
IN
       Unger, Evan C., Tucson, AZ, United States
       Shen, DeKang, Tucson, AZ, United States
Imarx Therapeutics, Inc., Tucson, AZ, United States (U.S. corporation)
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B1 20020903

20000203 (9)

PA

PΙ

ΑI

US 6444660

US 2000-496761

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Division of Ser. No. US 1997-851780, filed on 6 May 1997, now patented,
RLI
       Pat. No. US 6090800
       Utility
DT
       GRANTED
FS
       Primary Examiner: Badio, Barbara P.
EXNAM
       Woodcock Washburn LLP
LREP
       Number of Claims: 13
CLMN
       Exemplary Claim: 1
ECL
       0 Drawing Figure(s); 0 Drawing Page(s)
LN.CNT 6452
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention is directed to novel lipid soluble steroid
       prodrugs, compositions comprising steroid prodrugs, and uses of the
       same.
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 3 OF 24 USPATFULL
L7
       2002:217089 USPATFULL
AN
       Methods of using polynucleotide compositions
ΤI
       Kabanov, Alexander V., Omaha, NE, United States Alakov, Valery Y., Montreal, CANADA
IN
       Vinogradov, Serguie, Omaha, NE, United States
       Supratek Pharma Inc., CANADA (non-U.S. corporation)
PA
                                20020827
PΙ
       US 6440743
                          B1
       US 1999-320640
                                19990526 (9)
ΑT
       Division of Ser. No. US 1998-124943, filed on 30 Jul 1998, now patented,
RLI
       Pat. No. US 6221959 Continuation-in-part of Ser. No. US 1997-912968,
       filed on 1 Aug 1997, now patented, Pat. No. US 6353055
       Continuation-in-part of Ser. No. US 1994-342209, filed on 18 Nov 1994,
       now patented, Pat. No. US 5656611
DT
       Utility
FS
       GRANTED
EXNAM
       Primary Examiner: McGarry, Sean; Assistant Examiner: Epps, Janet
       Mathews, Collins, Shepherd & McKay, P.A. Number of Claims: 13
LREP
CLMN
ECL
       Exemplary Claim: 1
DRWN
       0 Drawing Figure(s); 0 Drawing Page(s)
LN.CNT 2206
CAS INDEXING IS AVAILABLE FOR THIS PATENT. .
       Compositions for stabilizing polynucleic acids and increasing the
       ability of polynucleic acids to cross cell membranes and act in the
       interior of a cell. In one aspect, the invention provides a
       polynucleotide complex between a polynucleotide and certain polyether
       block copolymers. The polynucleotide complex can further include a
       polycationic polymer, as well as suitable targeting molecules and
       surfactants. The invention also provides a polynucleotide complex
       between a polynucleotide and a block copolymer comprising a polyether
       block and a polycation block.
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 4 OF 24 USPATFULL
       2002:167866 USPATFULL
AN
ΤI
       Acoustically active drug delivery systems
IN
       Unger, Evan C., Tucson, AZ, United States
       Bristol-Myers Squibb Medical Imaging, Inc., Princeton, NJ, United States
PA
       (U.S. corporation)
PΙ
       US 6416740
                                20020709
       US 1998-75343
ΑI
                                19980511 (9)
PRAI
       US 1997-46379P
                           19970513 (60)
DT
       Utility
FS
       GRANTED
      Primary Examiner: Dudash, Diana; Assistant Examiner: Sharareh, Shahnam
EXNAM
LREP
       Woodcock Washburn LLP
```

CLMN Number of Claims: 15 ECL Exemplary Claim: 1

DRWN 9 Drawing Figure(s); 9 Drawing Page(s)

LN.CNT 5660

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The present invention is directed to targeted therapeutic delivery systems comprising a gas or gaseous precursor filled microsphere wherein said gas or gaseous precursor filled microsphere comprises an oil, a surfactant, and a therapeutic compound. Methods of preparing the targeted therapeutic delivery systems are also embodied by the present invention which comprise processing a solution comprising an oil and a surfactant in the presence of a gaseous precursor, at a temperature below the gel to liquid crystalline phase transition temperature of the surfactant to form gas or gaseous precursor filled microsphere, and adding to said microspheres a therapeutic compound resulting in a targeted therapeutic delivery system, wherein said processing is selected from the group consisting of controlled agitation, controlled drying, and a combination thereof.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 5 OF 24 USPATFULL

AN 2002:72457 USPATFULL

TI SOLID POROUS MATRICES AND METHODS OF MAKING AND USING THE SAME

UNGER, EVAN C., TUCSON, AZ, UNITED STATES

PI US 2002039594 A1 20020404 AI US 1998-75477 A1 19980511 (9)

PRAI US 1997-46379P 19970513 (60)

DT Utility

IN

FS APPLICATION

LREP WOODCOCK WASHBURN KURTZ, MACKIEWICZ AND NORRIS, ONE LIBERTY PLACE 46TH FLOOR, PHILADELPHIA, PA, 19103

CLMN Number of Claims: 106 ECL Exemplary Claim: 1

DRWN 1 Drawing Page(s)

LN.CNT 5207

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The present invention is directed to a solid porous matrix comprising a solvent and a surfactant in combination with a bioactive agent. The solvent and the surfactant may, if desired, form vesicles, an agglomeration of which comprises the matrix. The composition optionally comprises a gas or a gaseous precursor. The emulsion may be dried, and subsequently reconstituted in an aqueous or organic solution.

The present invention is also directed to a method of preparing a solid porous matrix comprising combining a solvent, a surfactant, and a therapeutic to form an emulsion; and processing the emulsion by controlled drying, or controlled agitation and controlled drying to form a solid porous matrix. The resulting solid porous matrix may also comprise a gas or gaseous precursor and be added to a resuspending medium.

A method for the controlled delivery of a targeted therapeutic to a region of a patient is another embodiment of the present invention. The method comprises administering to the patient a composition having a solid porous matrix comprising a solvent, a surfactant, a therapeutic, and a gas or gaseous precursor, monitoring the composition using energy to determine the presence of the composition in the region; and releasing the therapeutic from the composition in the region using energy.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 6 OF 24 USPATFULL AN 2002:57879 USPATFULL

```
Polynucleotide compositions for intramuscular administration
ΤI
       Lemieux, Pierre M., Ste.-Therese, CANADA
IN
       Kabanov, Alexander V., Omaha, NE, United States
       Alakov, Valery Y., D'Urfe, CANADA
       Vinogradov, Sergey V., Omaha, NE, United States
Supratek Pharma Inc., Doryal, United States (non-U.S. corporation)
PA
                                 20020319
                           B1
ΡI
       US 6359054
                                 19990108 (9)
       US 1999-227364
ΑI
       Continuation-in-part of Ser. No. US 1998-124943, filed on 30 Jul 1998,
RLI.
       now patented, Pat. No. US 6221959 Continuation-in-part of Ser. No. US
       1997-912968, filed on 1 Aug 1997 Continuation-in-part of Ser. No. US
       1994-342209, filed on 18 Nov 1994, now patented, Pat. No. US 5656611
DT
       Utility
FS
       GRANTED
       Primary Examiner: Szekely, Peter
EXNAM
       Mathews, Collins, Shepherd & Gould, P.A.
       Number of Claims: 25
CLMN
       Exemplary Claim: 1
ECL
       0 Drawing Figure(s); 0 Drawing Page(s)
DRWN
LN.CNT 2493
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Compositions and methods for intramuscular administration of
       polynucleotides, such as RNA, DNA, or derivatives thereof comprising
       polynucleotides and block copolymers of alkylethers. The invention also
       provides compositions and methods for stabilizing polynucleic acids and
       increasing the ability of polynucleic acids to cross cell membranes and
       act in the interior of a cell.
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 7 OF 24 USPATFULL
       2002:45673 USPATFULL
AN
ТT
       Polynucleotide compositions
IN
       Kabanov, Alexander Victorovich, Omaha, NE, United States
       Alakov, Valery Yulievich, D'Urfe, CANADA
Vingogradov, Sergey V., Omaha, NE, United States
       Supratek Pharma Inc., Quebec, CANADA (non-U.S. corporation)
PA
PΙ
       US 6353055
                           Bl
                                20020305
AΙ
       US 1997-912968
                                19970801 (8)
       Continuation-in-part of Ser. No. US 1994-342209, filed on 18 Nov 1994,
RLI
       now patented, Pat. No. US 5656611
DT
       Utility
FS
       GRANTED
EXNAM
       Primary Examiner: Szekely, Peter
       Mathews, Collins, Shepherd & Gould, P.A.
LREP
CLMN
       Number of Claims: 11
ECL
       Exemplary Claim: 1
DRWN
       0 Drawing Figure(s); 0 Drawing Page(s)
LN.CNT 2021
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The invention provides compositions for stabilizing polynucleic acids
       and increasing the ability of polynucleic acids to cross cell membranes
       and act in the interior of a cell. In one aspect, the invention provides
       a polynucleotide complex between a polynucleotide and certain polyether
       block copolymers. Preferably, the polynucleotide complex will further
       include a polycationic polymer. The compositions can further include
       suitable targeting molecules and surfactants. In another aspect, the
       invention provides a polynucleotide complex between a polynucleotide and
       a block copolymer comprising a polyether block and a polycation block.
       In yet another aspect, the invention provides polynucleotides 10 that
      have been covalently modified at their 5' or 3' end to attach a
      polyether polymer segment.
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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ANSWER 8 OF 24 USPATFULL
L.7
       2002:37868 USPATFULL
AN
       Methods and compositions for sealing tissue leaks
TT
       Wilkie, James, Melrose, MA, UNITED STATES
IN
       Rolke, James, Fitzwilliam, NH, UNITED STATES
       Burzio, Luis, Andover, MA, UNITED STATES
       Tammishetti, Shekharam, Secunderabad, INDIA
       Pendharkar, Sanyog Manohar, Oldbridge, NJ, UNITED STATES
                          A1
                               20020221
       US 2002022588
PΙ
       US 2000-747293
                          A1
                               20001222 (9)
AΙ
       Continuation-in-part of Ser. No. WO 1999-US14232, filed on 23 Jun 1999,
RLI
       UNKNOWN
       US 1998-90609P
                           19980623 (60)
PRAI
                           20000425 (60)
       US 2000-199469P
       US 1999-171859P
                           19991222 (60)
       Utility
DT
FS
       APPLICATION
       TESTA, HURWITZ & THIBEAULT, LLP, HIGH STREET TOWER, 125 HIGH STREET,
LREP
       BOSTON, MA, 02110
       Number of Claims: 167
CLMN
       Exemplary Claim: 1
ECL
DRWN
       No Drawings
LN.CNT 2885
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The invention provides methods and compositions that are useful for
       adhering biological and/or synthetic tissues, sealing fluid and/or
       gaseous leaks in biological and/or synthetic tissues, and preparing
       implants useful for delivery of a bioactive molecule such as a drug, for
       bulking applications, or for tissue prostheses. The present invention
       also relates to bio-erodable adhesive or occluding compositions and
       methods of using the same.
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 9 OF 24 USPATFULL
1.7
       2002:22133 USPATFULL
AN
       Novel drosophila tumor necrosis factor class molecule ("DmTNF") and
TI
       variants thereof
       Carroll, Pamela M., Princeton, NJ, UNITED STATES
IN
       Chen, Jian, Princeton, NJ, UNITED STATES
       Ramanathan, Chandra S., Wallingford, CT, UNITED STATES
       Xiao, Hong, Princeton Junction, NJ, UNITED STATES
       Guan, Bo, Princeton, NJ, UNITED STATES
       Bowen, Michael A., Lawrenceville, NJ, UNITED STATES
                          A1
                               20020131
PΤ
       US 2002012968
                               20010320 (9)
ΑI
       US 2001-813329
                          A1
PRAI
       US 2000-190816P
                           20000321 (60)
DT
      Utility
FS
      APPLICATION
      MARLA J MATHIAS, BRISTOL-MYERS SQUIBB COMPANY, PATENT DEPARTMENT, P O
LREP
       BOX 4000, PRINCETON, NJ, 08543-4000
CLMN
      Number of Claims: 40
       Exemplary Claim: 1
ECI.
       18 Drawing Page(s)
DRWN
LN.CNT 9244
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention provides novel polynucleotides encoding Drosophila
       DmTNF polypeptides, fragments and homologs thereof. The present
       invention also is directed to novel polynucleotides encoding two
       Drosophila DmTNF variants, DmTNFv1 and DmTNFv2 polypeptides, fragments
       and homologs thereof. Also provided are vectors, host cells, antibodies,
      and recombinant and synthetic methods for producing said polypeptides.
      The invention further relates to screening methods for identifying
      agonists and antagonists of the polynucleotides and polypeptides of the
      present invention, in addition to methods of genetically modifying
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Drosophila or cultured cells to express or mis-express DmTNF, DmTNFv1, or DmTNFv2. The invention also relates to the use of such modified insects or cells to characterize DmTNF activity, identify TNF-like genes and/or genes implicated in modulating TNF, characterize TNF signaling pathways, and/or to identify modulators of DmTNF activity.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

```
1.7
     ANSWER 10 OF 24 USPATFULL
        2001:234992 USPATFULL
AN
        Nanoqel networks and biological agent compositions thereof
ΤI
        Kabanov, Alexander V., Omaha, NE, United States Vinogradov, Sergey V., Omaha, NE, United States
TN
        Supratek Pharma, Inc., Canada (non-U.S. corporation)
PA
                                  20011225
                            B1
        US 6333051
PΤ
ΑI
        US 1998-146651
                                  19980903 (9)
        Utility
DT
        GRANTED
FS
EXNAM Primary Examiner: Riley, Jezia
       Mathews, Collins, Shepherd & Gould, P.A. Number of Claims: 12
LREP
CLMN
       Exemplary Claim: 1
ECL
DRWN
       No Drawings
LN.CNT 2246
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Copolymer networks having at least one cross-linked polyamine
       polymer fragment and at least one nonionic water-soluble polymer
        fragment, and compositions thereof, having at least one suitable
       biological agent.
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
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ANSWER 11 OF 24 USPATFULL
1.7
       2001:182086 USPATFULL
AN
TI
       Novel methods of ultrasound treatment using gas or gaseous
       precursor-filled compositions
TN '
       Unger, Evan C., Tucson, AZ, United States
PA
       ImaRx Pharmaceutical Corp. (U.S. corporation)
                               20011018
       US 2001031243
PΙ
                          A1
       US 2001-813484
                               20010321 (9)
ΑI
                          A1
       Division of Ser. No. US 1997-929847, filed on 15 Sep 1997, PENDING
RLI
DT
       Utility
       APPLICATION
FS
LREP
       Woodcock Washburn Kurtz, Mackiewicz & Norris LLP, 46th Floor, One
       Liberty Place, Philadelphia, PA, 19103
CLMN
       Number of Claims: 34
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
LN.CNT 6360
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention describes, among other things, the surprising
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The present invention describes, among other things, the surprising discovery that gaseous precursor filled compositions are profoundly more effective as acoustically active contrast agents when they are thermally preactivated to temperatures at or above the boiling point of the instilled gaseous precursor prior to their in vivo administration to a patient. Further optimization of contrast enhancement is achieved by administering the gaseous precursor filled compositions to a patient as an infusion. Enhanced effectiveness is also achieved for ultrasound mediated targeting and drug delivery.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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L7 ANSWER 12 OF 24 USPATFULL AN 2001:167740 USPATFULL
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TI Composition for treating benign prostatic hypertrophy

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Gokcen, Muharrem, Minneapolis, MN, United States
IN
       Guy, Terry J., Chaska, MN, United States
       Immunolytics, Inc., Minneapolis, MN, United States (U.S. corporation)
PA
                                20011002
       US 6296847
                          Bl
PΤ
                                19931117 (8)-
       US 1993-154158
ΑI
       Continuation of Ser. No. US 1991-707662, filed on 30 May 1991, now
RLI
       abandoned Continuation of Ser. No. US 1989-429966, filed on 31 Oct 1989,
       now abandoned Continuation-in-part of Ser. No. US 1989-303809, filed on
       27 Jan. 1989, now abandoned
       Utility
DT
       GRANTED
FS
       Primary Examiner: Witz, Jean C.
EXNAM
       Merchant & Gould P.C.
LREP
       Number of Claims: 31
CLMN
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
LN.CNT 3351
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The invention provides a composition and method for treating benign
       prostatic hypertrophy in mammals so as to cause the dissolution and
       regression of hypertrophied prostatic tissue and thereby provide relief
       from the obstructive symptoms associated with the disease. The present
       composition preferably comprises a sterile pyrogen-free solution of the
       hydrolytic enzymes collagenase and hyaluronidase, a nonionic surfactant,
       and an antibiotic; all provided, in a pharmaceutically acceptable,
       buffered, isotonic, aqueous carrier. The present method preferably
       comprises the direct intraprostatic injection of a safe and
       therapeutically effective dose of the composition via the transurethral
       route of administration.
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 13 OF 24 USPATFULL
       2001:144937 USPATFULL
AN
ΤŦ
       Solid matrix therapeutic compositions
       Unger, Evan C., Tucson, AZ, United States
ΤN
       ImaRx Therapeutics, Inc. (U.S. corporation)
PA
PΙ
       US 2001018072
                          A1
                               20010830
       US 2001-828762
                               20010409 (9)
ΑT
                          Α1
RLI
       Division of Ser. No. US 1998-75477, filed on 11 May 1998, PENDING
PRAI
       US 1997-46379P
                           19970513 (60)
DT
       Utility
FŞ
       APPLICATION
       Mackiewicz & Norris LLP, One Liberty Place - 46th Floor, Philadelphia,
LREP
       PA, 19103
CLMN
       Number of Claims: 38
ECL
       Exemplary Claim: 1
DRWN
       1 Drawing Page(s)
LN.CNT 4899
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention is directed to a solid porous matrix comprising a
       surfactant in combination with a bioactive agent. The solid porous
       matrix may be prepared by combining a surfactant and a therapeutic,
       together with a solvent, to form an emulsion containing random
       aggregates of the surfactant and the therapeutic, and processing the
       emulsion by controlled drying, or controlled agitation and controlled
       drying to form the solid porous matrix.
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 14 OF 24 USPATFULL
AN
       2001:130897 USPATFULL
ΤI
       Prolonged release of GM-CSF
       Gombotz, Wayne R., Kirkland, WA, United States
       Pettit, Dean K., Seattle, WA, United States
```

Pankey, Susan C., Yardley, PA, United States Immunex Corporation, Seattle, WA, United States (U.S. corporation) PA US 6274175 20010814 PΤ В1 19991117 (9) US 1999-442370 ΑI Continuation of Ser. No. US 1998-185213, filed on 3 Nov 1998, now RLI patented, Pat. No. US 6120807 Division of Ser. No. US 1995-542445, filed on 12 Oct 1995, now patented, Pat. No. US 5942253 DT GRANTED FS EXNAM Primary Examiner: Azpuru, Carlos A Sheiness, Diana K. LREP Number of Claims: 24 CLMN Exemplary Claim: 1 ECL 11 Drawing Figure(s); 6 Drawing Page(s) DRWN LN.CNT 1524 CAS INDEXING IS AVAILABLE FOR THIS PATENT. Formulations for controlled, prolonged release of GM-CSF have been AB developed. These are based on solid microparticles formed of the combination of biodegradable, synthetic polymers such as poly(lactic acid) (PLA), poly(glycolic acid) (PGA), and copolymers thereof with excipients and drug loadings that yield zero order or first order release, or multiphasic release over a period of approximately three to twenty one days, preferably one week, when administered by injection. In the preferred embodiment, the microparticles are microspheres having diameters in the range of 10 to 60 microns, formed of a blend of PLGA having different molecular weights, most preferably 6,000, 30,000 and 41,000. Other embodiments have been developed to alter the release kinetics or the manner in which the drug is distributed in vivo. For example, in some cases a polymer is selected which elicits a mild inflammatory reaction, for example, PLGA and polyanhydrides can act as chemoattractant, either due to the polymer itself or minor contaminants in the polymer, or polymers which are bloadhesive are used for transmucosal or oral delivery. In another embodiment, the GM-CSF is administered in a hydrogel which can be injected subcutaneous or at a specific site for controlled release. The microparticles or hydrogel are administered to the patient in an amount effect to stimulate proliferation of hematopoietic cells, especially white cells. CAS INDEXING IS AVAILABLE FOR THIS PATENT. ANSWER 15 OF 24 USPATFULL AN 2001:97899 USPATFULL TΤ Autocross-linked hyaluronic acid and related pharmaceutical compositions for the treatment of arthropathies IN Bellini, Davide, Padua, Italy Paparella, Annamaria, Bari, Italy O'Regan, Michael, Padua, Italy Callegaro, Lanfranco, Vicenza, Italy Fidia, S.p.A., Abano Terme, Italy (non-U.S. corporation) PA PI US 6251876 20010626 B1 WO 9749412 19971231 US 1999-202817 19990625 (9) AΙ WO 1997-EP3238 19970620 PCT 371 date 19990625 19990625 PCT 102(e) date PRAI IT 1996-PD163 19960621 DT Utility FS GRANTED EXNAM Primary Examiner: Peselev, Elli LREP Birch, Stewart, Kolasch & Birch LLP, Svensson, Leonard R. CLMN Number of Claims: 8 ECL Exemplary Claim: 1,2,8 19 Drawing Figure(s); 17 Drawing Page(s)

LN.CNT 1233

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The present invention relates to compositions containing an autocross-linked form of hyaluronic acid as a first component in a mixture with a second component noncross-linked hyaluronic acid, and possibly also in combination with another pharmacologically active substance. These compositions can be used in the treatment of arthropathies due to their unique viscoelastic properties.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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ANSWER 16 OF 24 USPATFULL
1.7
        2001:59978 USPATFULL
AN
        Polynucleotide compositions
TI
        Kabanov, Alexander V., Omaha, NE, United States
IN
       Alakov, Valery Y., D'Urfe, Canada
       Vinogradov, Sergey V., Omaha, NE, United States
PΑ
       Supratek Pharma, Inc., Montreal, Canada (non-U.S. corporation)
                                  20010424
PI
       US 6221959
                            B1
       US 1998-124943 19980730 (9)
Continuation-in-part of Ser. No. US 1998-912968, filed on 1 Aug 1998
Continuation-in-part of Ser. No. US 1994-342209, filed on 18 Nov 1994,
ΑI
RLI
       now patented, Pat. No. US 5656611
DT
       Utility
FS
       Granted
EXNAM Primary Examiner: Michl, Paul R.
       Mathews, Collins, Shepherd & Gould, P.A.
LREP
       Number of Claims: 8
CLMN
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
LN.CNT 2309
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Compositions for stabilizing polynucleic acids and increasing the
       ability of polynucleic acids to cross cell membranes and act in the
       interior of a cell. In one aspect, the invention provides a
       polynucleotide complex between a polynucleotide and certain polyether
       block copolymers. The polynucleotide complex can further include a
       polycationic polymer, as well as suitable targeting molecules and
       surfactants. The invention also provides a polynucleotide complex
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.

block and a polycation block.

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ANSWER 17 OF 24 USPATFULL
ΑN
       2000:124586 USPATFULL
TΤ
       Prolonged release of GM-CSF
TN
       Gombotz, Wayne, Kirkland, WA, United States
       Pettit, Dean, Seattle, WA, United States
       Pankey, Susan, Seattle, WA, United States
       Immunex Corporation, Seattle, WA, United States (U.S. corporation)
PA
PΙ
       US 6120807
                                20000919
ΑI
       US 1998-185213
                                19981103 (9)
RLI
       Division of Ser. No. US 1995-542445, filed on 12 Oct 1995, now patented,
       Pat. No. US 5942253
DT
       Utility
FS
       Granted
EXNAM
       Primary Examiner: Azpuru, Carlos A.
       Arnall Golden & Gregory, LLP
LREP
CLMN
       Number of Claims: 23
ECL
       Exemplary Claim: 1
DRWN
       11 Drawing Figure(s); 6 Drawing Page(s)
LN.CNT 1382
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     . Formulations for controlled, prolonged release of GM-CSF have been
       developed. These are based on solid microparticles formed of the
       combination of biodegradable, synthetic polymers such as poly(lactic
```

between a polynucleotide and a block copolymer comprising a polyether

acid) (PLA), poly(glycolic acid) (PGA), and copolymers thereof with excipients and drug loadings that yield zero order or first order release, or multiphasic release over a period of approximately three to twenty one days, preferably one week, when administered by injection. In the preferred embodiment, the microparticles are microspheres having diameters in the range of 10 to 60 microns, formed of a blend of PLGA having different molecular weights, most preferably 6,000, 30,000 and 41,000. Other embodiments hare been developed to alter the release kinetics or the manner in which the drug is distributed in vivo. For example, in some cases a polymer is selected which elicits a mild inflammatory reaction, for example, PLGA and polyanhydrides can act as chemoattractant, either due to the polymer itself or minor contaminants in the polymer, or polymers which are bloadhesive are used for transmucosal or oral delivery. In another embodiment, the GM-CSF is administered in a hydrogel which can be injected subcutaneous or at a specific site for controlled release. The microparticles or hydrogel are administered to the patient in an amount effect to stimulate proliferation of hematopoietic cells, especially white cells.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

```
ANSWER 18 OF 24 USPATFULL
L7
       2000:91955 USPATFULL
ΑN
       Lipid soluble steroid prodrugs
ΤI
IN
       Unger, Evan C., Tucson, AZ, United States
       Shen, DeKang, Tucson, AZ, United States
       Imarx Pharmaceutical Corp., Tucson, AZ, United States (U.S. corporation)
PA
                               20000718
       US 6090800
ΡI
       US 1997-851780
                               19970506 (8)
AΙ
DT
       Utility
FS
       Granted
EXNAM Primary Examiner: Dees, Jose' G.; Assistant Examiner: Badio, Barbara
       Woodcock Washburn Kurtz Mackiewicz & Norris LLP
LREP
       Number of Claims: 10
CLMN
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
LN.CNT 6285
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention is directed to novel lipid soluble steroid
       prodrugs compositions comprising steroid prodrugs, and uses of the same.
```

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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ANSWER 19 OF 24 USPATFULL
1.7
       2000:87694 USPATFULL
AN
ТT
       Compositions of microspheres for wound healing
IN
       Ritter, Vladimir, Kiriat-Yam, Israel
       Ritter, Marina, Kiriat-Yam, Israel
PA
       Polyheal Ltd., Haifa, Israel (non-U.S. corporation)
PΙ
       US 6086863
                               20000711
                               19981023 (9)
ΑI
       US 1998-177954
       Continuation-in-part of Ser. No. US 1997-868950, filed on 4 Jun 1997,
RLI
       now patented, Pat. No. US 5861149
DT
       Utility
FS
       Granted
EXNAM
      Primary Examiner: Cintins, Marianne M.; Assistant Examiner: Kim, Vickie
LREP
       Graham & James LLP
CLMN
       Number of Claims: 31
ECL
       Exemplary Claim: 1
DRWN
       30 Drawing Figure(s); 30 Drawing Page(s)
LN.CNT 1659
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Therapeutic compositions of microspheres for application to wounds
       and/or lesions for accelerating wound healing and muscle regeneration.
       The microspheres are made up of non-biodegradable material having a
```

substantial surface charge. The therapeutic composition further includes a pharmaceutically acceptable carrier in which the microspheres are insoluble and a container for holding the composition. The therapeutic composition further contains pharmacologic agents or biologics that accelerate the wound healing process.

```
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 20 OF 24 USPATFULL
L7
        2000:21560 USPATFULL
AN
        Prodrugs comprising fluorinated amphiphiles
ΤI
        Unger, Evan C., Tucson, AZ, United States
IN
        Imarx Pharmaceutical Corp., Tucson, AZ, United States (U.S. corporation)
PA
                                  20000222
PΙ
        US 6028066
        US 1997-887215
                                  19970702 (8)
ΑI
       Continuation-in-part of Ser. No. US 1997-851780, filed on 6 May 1997
RLI
DT
       Utility
        Granted
FS
       Primary Examiner: Dees, Jose' G.; Assistant Examiner: Badio, Barbara
EXNAM
        Woodcock Washburn Kurtz Mackiewicz & Norris LLP
LREP
       Number of Claims: 8
CLMN
ECL
       Exemplary Claim: 1
DRWN
       No Drawings
LN.CNT 6329
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention describes, inter alia, novel prodrugs comprising
        fluorinated amphiphiles, compositions comprising the novel prodrugs, and
       methods of use of the prodrugs and compositions.
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
1.7
     ANSWER 21 OF 24 USPATFULL
       1999:99401 USPATFULL
AN
       Prolonged release of GM-CSF
TΤ
IN
       Gombotz, Wayne, Kirkland, WA, United States
       Pettit, Dean, Seattle, WA, United States
       Pankey, Susan, Seattle, WA, United States
       Lawter, James Ronald, Goshen, NY, United States
Huang, W. James, Sommerville, NJ, United States
Immunex Corporation, Seattle, WA, United States (U.S. corporation)
PA
       American Cyanamid Company, Pearl River, NY, United States (U.S.
       corporation)
PΤ
       US 5942253
                                 19990824
ΑI
       US 1995-542445
                                 19951012 (8)
DT
       Utility
FS
       Granted
EXNAM
       Primary Examiner: Azpuru, Carlos
LREP
       Arnall Golden & Gregory, LLP
       Number of Claims: 27
CLMN
ECL
       Exemplary Claim: 1
DRWN
       11 Drawing Figure(s); 6 Drawing Page(s)
LN.CNT 1403
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
```

Formulations for controlled, prolonged release of GM-CSF have been developed. These are based on solid microparticles formed of the combination of biodegradable, synthetic polymers such as poly(lactic acid) (PLA), poly(glycolic acid) (PGA), and copolymers thereof with excipients and drug loadings that yield zero order or first order release, or multiphasic release over a period of approximately three to twenty one days, preferably one week, when administered by injection. In the preferred embodiment, the microparticles are microspheres having diameters in the range of 10 to 60 microns, formed of a blend of PLGA having different molecular weights, most preferably 6,000, 30,000 and 41,000. Other embodiments have been developed to alter the release kinetics or the manner in which the drug is distributed in vivo. For

example, in some cases a polymer is selected which elicits a mild inflammatory reaction, for example, PLGA and polyanhydrides can act as chemoattractant, either due to the polymer itself or minor contaminants in the polymer, or polymers which are bioadhesive are used for transmucosal or oral delivery. In another embodiment, the GM-CSF is administered in a hydrogel which can be injected subcutaneous or at a specific site for controlled release. The microparticles or hydrogel are administered to the patient in an amount effect to stimulate proliferation of hematopoietic cells, especially white cells.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

mammals as formed objects or depots.

```
ANSWER 22 OF 24 USPATFULL
       1998:75722 USPATFULL
AN
       Products comprising substrates capable of enzymatic cross-linking
ТT
       Cappello, Joseph, San Diego, CA, United States
TN
       Protein Polymer Technologies, San Diego, CA, United States (U.S.
PA
       corporation)
PΙ
       US 5773577
                               19980630
                               19950302 (8)
       US 1995-397633
AΤ
       Continuation-in-part of Ser. No. US 1994-205518, filed on 3 Mar 1994,
RLI
      now abandoned
DT
       Utility
FS
       Granted
      Primary Examiner: Patterson, Jr., Charles L.; Assistant Examiner: Stole,
EXNAM
      Einar
       Trecartin, Richard F.Flehr Hohbach Test Albritton & Herbert LLP
LREP
      Number of Claims: 29
CLMN
      Exemplary Claim: 1
ECL
DRWN
      No Drawings
LN.CNT 3006
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      Polymers are provided comprising protein polymers comprising blocks of
      repeating units and sequences comprising amino acids, individually or in
      defined sequences, capable of enzyme catalyzed covalent bond formation
      for cross-linking, as exemplified by glutamine and/or lysine reactive
      for FXIII catalyzed isopeptide formation or non-amino acid polymers
      having side chains comprising such amino acids or sequences, which may
      be used for preparation of articles of manufacture, particularly
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cross-linkable compositions. By appropriate choice of the polymer,

resorbable implantable polymers may be used in internal applications for

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CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 23 OF 24 USPATFULL
       97:93905 USPATFULL
AN
ТT
       Crosslinked carboxy polysaccharides
IN
       Della Valle, Francesco, Padua, Italy
       Romeo, Aurelio, Rome, Italy
       Fidia, S.p.A., Abano Terme, Italy (non-U.S. corporation)
PA
       US 5676964
                               19971014
PI
       US 1995-465055
ΑI
                               19950605 (8)
       Continuation of Ser. No. US 1993-70505, filed on 1 Jun 1993 which is a
RLI
       continuation of Ser. No. US 1989-350919, filed on 12 May 1989, now
       abandoned
       IT 1988-47964
                         . 19880513
PRAI
DT
       Utility
FS
       Granted
EXNAM
      Primary Examiner: Peselev, Elli
      Birch, Stewart, Kolasch & Birch, LLP
LREP
CLMN
      Number of Claims: 65
      Exemplary Claim: 1,36
ECL
DRWN
      No Drawings
LN.CNT 2523
```

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CAS INDEXING IS AVAILABLE FOR THIS PATENT.
        Inter and/or intramolecular cross-linked esters of acid
AB
        polysaccharides are disclosed in which a part or all of the
        carboxy groups are esterified with hydroxyl groups of the same
        molecule and/or of different molecules of the acid
        polysaccharide. These inner cross-linked esters of
        polysaccharide acids are useful in the field of biodegradable
        plastic materials, to manufacture sanitary and surgical articles, in the
        cosmetic and pharmaceutical fields, in the food industry and in many
        other industrial fields.
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 24 OF 24 USPATFULL
        92:42541 USPATFULL
AN
        Method for treating benign prostatic hypertrophy
TΤ
        Gokcen, Muharrem, Minneapolis, MN, United States
IN
        Guy, Terry J., Chaska, MN, United States
Immunolytics, Inc., Minneapolis, MN, United States (U.S. corporation)
PA
        US 5116615
                                 19920526
PT
        us 1991-707628
                                 19910530 (7)
AΙ
        Continuation of Ser. No. US 1989-429966, filed on 31 Oct 1989, now
        abandoned which is a continuation-in-part of Ser. No. US 1989-303809,
        filed on 27 Jan 1989, now abandoned
DT
        Utility
FS
        Granted
EXNAM Primary Examiner: Stone, Jacqueline
LREP
       Merchant, Gould, Smith, Edell, Welter & Schmidt
CLMN
       Number of Claims: 19
        Exemplary Claim: 1
ECL
DRWN
       No Drawings
LN.CNT 3209
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The invention provides a composition and method for treating benign
AB
       prostatic hypertropy in mammals so as to cause the dissolution and
       regression of hypertrophied prostatic tissue and thereby provide relief
        from the obstructive symptoms associated with the disease. The present
       composition preferably comprises a sterile pyrogen-free solution of the
       hydrolytic enzymes collagenase and hyaluronidase, a nonionic surfactant,
        and an antibiotic; all provided, in a pharmaceutically acceptable,
       buffered, isotonic, aqueous carrier. The present method preferably
       comprises the direct intraprostatic injection of a safe and
       therapeutically effective dose of the composition via the transurethral
       route of administration.
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
=> dis hist
     (FILE 'HOME' ENTERED AT 14:30:46 ON 23 SEP 2002)
     FILE 'BABS, CAPLUS, CBNB, CEN, CIN, DKILIT, IFIPAT, JICST-EPLUS, PASCAL, PLASNEWS, PROMT, RAPRA, SCISEARCH, TEXTILETECH, USPATFULL, USPAT2,
     WPINDEX, WTEXTILES' ENTERED AT 14:31:14 ON 23 SEP 2002
Ll
         218150 S POLYSACCHARIDE
L2
           7277 S L1 AND CARBOXY
L3
           3796 S L2 AND ACTIVAT?
            317 S L3 AND CROSS-LINK
L5
             76 S L4 AND (DIAMINE OR POLYAMINE)
L6
              0 S L5 AND HYALURONI
             24 S L5 AND HYALURONIC
=> s 17 and sulfation
```

1 L7 AND SULFATION

=> dis 18 bib abs

ANSWER 1 OF 1 USPATFULL $\Gamma8$ 2002:22133 USPATFULL AN Novel drosophila tumor necrosis factor class molecule ("DmTNF") and TΤ variants thereof Carroll, Pamela M., Princeton, NJ, UNITED STATES IN Chen, Jian, Princeton, NJ, UNITED STATES Ramanathan, Chandra S., Wallingford, CT, UNITED STATES Xiao, Hong, Princeton Junction, NJ, UNITED STATES Guan, Bo, Princeton, NJ, UNITED STATES Bowen, Michael A., Lawrenceville, NJ, UNITED STATES US 2002012968 A1 20020131 PΤ US 2001-813329 A1 20010320 (9) ΑI US 2000-190816P 20000321 (60) PRAI DT Utility APPLICATION FS MARLA J MATHIAS, BRISTOL-MYERS SQUIBB COMPANY, PATENT DEPARTMENT, P O LREP BOX 4000, PRINCETON, NJ, 08543-4000 Number of Claims: 40 CLMN ECL Exemplary Claim: 1 18 Drawing Page(s) DRWN LN.CNT 9244 CAS INDEXING IS AVAILABLE FOR THIS PATENT. The present invention provides novel polynucleotides encoding Drosophila DmTNF polypeptides, fragments and homologs thereof. The present invention also is directed to novel polynucleotides encoding two Drosophila DmTNF variants, DmTNFv1 and DmTNFv2 polypeptides, fragments and homologs thereof. Also provided are vectors, host cells, antibodies, and recombinant and synthetic methods for producing said polypeptides. The invention further relates to screening methods for identifying agonists and antagonists of the polynucleotides and polypeptides of the present invention, in addition to methods of genetically modifying Drosophila or cultured cells to express or mis-express DmTNF, DmTNFv1, or DmTNFv2. The invention also relates to the use of such modified

insects or cells to characterize DmTNF activity, identify TNF-like genes and/or genes implicated in modulating TNF, characterize TNF signaling

pathways, and/or to identify modulators of DmTNF activity.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

=> file chemistry
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FULL ESTIMATED COST

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FILE 'WSCA' ENTERED AT 14:36:29 ON 23 SEP 2002 COPYRIGHT (C) 2002 PAINT RESEARCH => s polysaccharide 239171 POLYSACCHARIDE L9 => s 19 and carboxy 628 L9 AND CARBOXY => s 110 and activat? 26 FILES SEARCHED... 40 L10 AND ACTIVAT? L11 => s lll and cross-link? 25 FILES SEARCHED... 1 L11 AND CROSS-LINK? L12 => dis 112 bib abs L12 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS 1990:406740 CAPLUS AN DN 113:6740 TΙ Preparation of crosslinked carboxy polysaccharides as biodegradable plastic materials for cosmetics and pharmaceuticals Della Valle, Francesco; Romeo, Aurelio IN PA Fidia S.p.A., Italy Eur. Pat. Appl., 37 pp. SO CODEN: EPXXDW DT Patent English FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE . --------------------EP 1989-108630 19890512 PΙ EP 341745 A1 19891115 EP 341745 19941214 В1 R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE WO 1989-EP519 19890512 WO 8910941 A1 19891116 W: AU, DK, FI, HU, JP, KR 19891129 AU 1989-35747 19890512 AU 8935747 A1 AU 631125 B2 19921119 HU 53666 A2 19901128 HU 1989-3636 19890512 HU 210926 В 19950928 19901129 T2 JP 1989-505458 19890512 JP 02504163 JP 2941324 B2 19990825 A2 19940914 EP 1994-108633 19890512 EP 614914 EP 614914 A3 19941228 EP 614914 В1 20000816 R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE ES 2064378 Т3 19950201 ES 1989-108630 19890512 IL 1989-90274 IL 90274 A1 19960912 19890512 CA 1339122 A1 19970729 CA 1989-599557 19890512 JP 10324701 A2 19981208 JP 1998-152832 19890512 AT 195534 E 20000915 AT 1994-108633 19890512 ES 2151910 Т3 20010116 ES 1994-108633 19890512 DK 9000109 DK 1990-109 19900312 19900112 Α US 5676964 Α 19971014 US 1995-465055 19950605 PRAI IT 1988-47964 Α 19880513 EP 1989-108630 А3 19890512 JP 1989-505458 **A3** 19890512 US 1989-350919 В1 19890512 WO 1989-EP519 Α 19890512 US 1993-70505 A1 19930601 AB Inter- and/or intramol. esters of acid polysaccharides contg.

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carboxy functions (e.g. auto-crosslinked polysaccharides), wherein (1) a first portion or all of the carboxy groups are esterified with hydroxy groups of the same mol. and/or of different mols. of the acid polysaccharide and/or (2) a second portion of the carboxy groups are esterified with a mono- or polyvalent alcs. including various drugs (e.g. alkaloids, anesthetic, analgesic, antiinflammatory, antiviral, antibacterial, etc.) and optionally salified with an alkali or alk. earth metal, Mg, Al, or an amine including various drugs (e.g. alkaloids, peptides, antipsychotics, phenothiazine, vasoconstrictors, etc.), are prepd. by treating an acidic polysaccharide (e.g., hyaluronic acid, alginic acid, CM-cellulose, carboxymethylchitin) with an activating agent (e.g. 2-chloro-1-methylpyridinium iodide) and subjecting the resulting intermediate activated polysaccharide derivs. to heat or irradn. These auto-crosslinked polysaccharide acids are useful in the field of biodegradable plastic materials to manuf. sanitary and surgical articles (e.g. surgical suture thread, film for artificial skin, and sponges for the medication of wounds and lesions), for pharmaceutical vehicles for controlled-release of drugs (capsules for the s.c. implantation of medicaments or microcapsules for s.c., i.m., or i.v. injection), etc.

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---Logging off of STN---

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	. ENTRY	SESSION
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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
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